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TITLE: Method and apparatus for detecting hydrogen cyanide

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Patented Dec. 19, 1950 2)534,229 @@UNITE'D STATES PAT,ENT @OFFICE 2,534,229 METHOD-AND-APPARATUS@FORIDETEC,T)ING. @HYDROGEN CYANIDE 'Homer W. Citrhart, and John A. Krynitsk @ Y, , Washington, D. C. Applicatioti'March'. 15,1945, Serial No. 582,965 15- Cl@tiins. (Cl. 23-232), (Granted under the act of - Mareh 4 3, 1883@ as amended 7@April 30, 1928, 1 370 0.@ G@ 757) Our invention relates 1:@tol. the @ detection, of 'hYr@ dr-ogen- eya@nide or hydrocyanic, acid in; admixture with air or other gases: @ U-nder ordinary conditions, hydrocyanic . or prussic-@acid. i@-a light gas haN7ing-an,@iodor@ re- 5 sembling:that:ofbitter-a-Imonds@ One lof@its.,best know-n properties @ is@ @its@ extreme toxicity @ and- for this -reason it is frequently- used@ as.@:a@ fumigat- ing agent. Another very common industrial ap- plication of cyanide, com@pounds is in electro-plat- 10 ing,-work where, solutions containing, the cyanide radical, are-conventional. The slig ht-hydrolysis and,,decomposition of @.these-.compounds irl solu- tion,,results in.thecontamination.of the.@a-tmos.. pher.e,,in.-.electro@-plating. room wit h-appreciable 15 concentrations - In, @addition to. the. i@ndustritil. uses, of cyanido com 6 unds, hydrocyanic@ acid itself. has-considbr. p able value@,as.,a.,war-,gas. In.spite of it,s volatility and, hence, its, lack,of persistency, -it is, very ef@. 20 fective because of its. extreme, toxi, city and lack of Isuch, aetion, as 1: @chrymation or., irritation. It is.7a.well-establishe.d fact..that 'many people can-not. detect the odor- of ..hydrocyanic acid. even when @it is present -in.. the, atmosphere in concen- 25 trations, far.. exceeding,, lethal.. concentrations. It, is, an object, of .. our .. invention. to, provide a simple and .convenient method . of ., detecting - containination of the-atmosphere.or other gases -with trae,es- of . hyd@tocyanic acid in amounts below @,,0 lethal 'concentrations. It is a second object of our invention to provide a device which can -be used, simply and easils, to detect small concentrations of hydrocyanic acid in small'samples@@of,air @or@other,@gas. 35 It is @another. @bbiect@of -@our-,Invention..to provide a @@ colorimetricquz@littctive;; and @ Bemi@ quantita; tive test apparatus- and: a @ composition.l.fori the.i. detee- tion. 6f., hydr@ocyanic - acid @ ound estimation of, @@ @it8 concentration; in a, contaminated@atmosphere. 40 A: further object of @.our. inventionl, is, to @provide a. method -@of @ preserving @,indefinitely chemica;l@. re" agents which are subject.@.to@%@decomi>osition' by contact@ with oxygen. @ Other objects and @ advantages. Of our invention 45 will, in, part, be @ obvious. and,@in:@part@ appear,here- inafter' I Our invention@ comprises the, @composition, methodl,and ' apparatus for preserving an oxygensensiti, ve@chemical@and,:for@detecting.-hydrocyanic 50 acid%bya@sharp colorimetric reaction.@.w.-hich will be described @in dettbil,and illustrated @@iir the drawing.; accompanying. this-. specification. @The droiiing@represents@,a@@longitudinal,@eross.se6tion.through- a; Itube comprisiig.: @the appara-tus 55 of.".our. @invention., A satisfactory colorimetric reaction must be one which is, first; extremely sensitive, second, one which gives a very@ sharp color change which is readily distinguished, and, third" one whir, h Preferably takes iplace, -Gn, a,, white background. There are a number of reactions of hydrocyanic acid which-will -give colbr, among which is the reaction@of hydrocyanic@acid,with alkaline sodium Picratetafbrmisopurpuricacid. Asatestforthe detection of hydrocyanic acid, this reaction is not P articularly, satisf actory for the color change @involved is one in which yellow- sodium pierate is converted to a reddish--brown iso-purpuric, acid. I-lydrocyanic. acid@ can also be made to react with cupric sulfide. However, - the reaction is not a good colorimetric one for it @ involies the con" version of 'the@ bia-ek cupric -sulfide to; f orm.white cuprous- cyanide The-reaction, of the cyanide radica; l@or hydrocyanic, acid with iron salts to-give ferri- ferrocyanide is,-very well known. I-lowever, it is to 0 complicated for convenient field use. We-have discovered@ o@ method whereby@ the reaction of-hydroeyonic acid with a cupric saitorganic aromatic, amine@ reagent can be used for the@ qualitative,.and: semi-quantitative detection of. hydrocyanic acid. The reaction gives@ a sharp color change and possesses the, advantage of being, rapid, direct and quite specific for hydroeyanic acid. In the past the principal obstacle to@.the use.. of

this type of- reaction,, that is, the cupric- salt-organic aromatic amine reagent, for the detection of hydrocyanic acid was the instability. of the various formulations. Short periods of '@ exposure to traces - of oxygen discolored the material so badly it was made useless for,col.orirnetriework@. We have discovered that a cupric salt-aroinatic-amine@combination which is highly sensi'Live to hydrocyanic acid and, consequently, can be used@ to detect micro- concentrations of hydroeyanic acid, can be stabilized and preserved so that the rea.@ent once prepared P,,nd se@iled into the apparatu-sr will remain %in operative condition for an indefinite period. In general we have found that organic aromatic; amines: such as @ ariiiine, ortho- phenylenediamine, para-toluidine, para-phenetidine, diphenylamine, dimethylaniline, N-@benzylaniline, phenyl-alphanaphthyl :amine, -bis.@(p-methylaminophenyl) methane, Michler's @ ketone, dibenzylaniline, 4,41 -@ tetramethyldiaminotriphenyl:methane, @414! 14"- hexamethyltriaminotriphenylmethane, p@.@di ethylaminodiphenyjmethane, dipheqylbenzidine, and tetramethyl diaminodiphenylinethane-, (or@ bis-@p- dimeth aminophenyl). Yi